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By:

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Patent

Attorney's Docket No. 028870-178

**IN THE UNITED STATES PATENT AND TRADEMARK OFFICE
UTILITY PATENT
APPLICATION TRANSMITTAL LETTER**

Box PATENT APPLICATION

Assistant Commissioner for Patents
Washington, D.C. 20231

Sir:

Enclosed for filing is the utility patent application of David C. Greenspan, Sean Lee and Marlo tan Walpole for Anti-Inflammatory Bioactive Glass Particulates.

Also enclosed are:

- [X] 8 sheet(s) of [] formal [X] informal drawing(s);
- [X] To follow: verified statement(s) claiming small entity status;
- [X] An [] executed [X] unexecuted declaration of the inventor(s)
[X] also is enclosed [] will follow.
- [X] Please amend the specification by inserting before the first line the sentence --This application claims priority under 35 U.S.C. §§ 119 and/or 365 to Provisional Application No. 60/131,529 filed on April 29, 1999, which designates the United States.--the entire content of which is hereby incorporated by reference.

CLAIMS					
	NO. OF CLAIMS		EXTRA CLAIMS	RATE	FEE
Basic Application Fee				(101)	\$ 690.00
Total Claims	11	MINUS 20 =	-0-	x \$18 = (103)	-0-
Independent Claims	4	MINUS 3 =	1	x \$78 = (102)	78.00
If multiple dependent claims are presented, add \$260.00 (104)					-0-
Total Application Fee					768.00
If verified statement claiming small entity status is enclosed, subtract 50% of Total Application Fee					384.00
Add Assignment Recording Fee of \$40.00 (581) if Assignment document is enclosed					-0-
TOTAL APPLICATION FEE DUE (Fee not enclosed)					\$384.00

**21839**

09560475, 042800

[X] This application is being filed without a filing fee. Issuance of a Notice to File Missing Parts of Application is respectfully requested.

This paper is submitted in duplicate.

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ANTI-INFLAMMATORY BIOACTIVE GLASS PARTICULATES

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ANTI-INFLAMMATORY BIOACTIVE GLASS PARTICULATES

FIELD OF THE INVENTION

5 The present invention relates to compositions and methods for use in the transient suppression of the inflammatory response by suppressing plasma levels of tissue necrosis factor-alpha and increasing plasma levels of interleukin-6.

BACKGROUND OF THE INVENTION

10 When an injury occurs, cell damage initially comes from the precipitating event, such as a cut, resulting in ruptured cells and severed or crushed capillaries and other blood vessels. The interruption of blood flow results in anoxia, causing the death of additional cells. The wound site quickly fills with dead and dying cells, extracellular substances (collagen,
15 elastic fibers, fat and ground substances), extravasated blood, and possibly bacteria and viruses introduced during the injury.

Tissue damage is not restricted to the initial area of injury, and may increase over the next several hours or days as a result of the release of
20 lysosomal enzymes from the injured cells or as a consequence of inflammation (swelling) and/or infection. (See Reese et al., Role of Fibronectin in Wound Healing, the subject matter of which is hereby incorporated by reference.) The inflammatory response is one of the normal stages of wound healing, and is necessary for subsequent phases of healing.

25 Inflammation is a vital process necessary for an organism to survive an external insult, such as a wound or burn. However, if unchecked, inflammation can have harmful consequences. For example, many chronic and even life-threatening disorders, such as asthma, rheumatoid arthritis,
30 lung fibrosis, peritoneal adhesions, hypersensitivity and autoimmune

diseases are a result of an uncontrolled inflammatory response. An unresolved inflammation in the lung resulting from bacterial infection (i.e., pneumonia) may eventually lead to extensive tissue damage and a chronic lung abscess. Inflammation of the peritoneal cavity, for example, and the resulting adhesions following abdominal surgery is a major cause of infertility in women. Asthma is an often life-threatening disorder which results from an inadvertently stimulated inflammatory response in the lungs. An excessive inflammatory response can cause extensive swelling, which can lead to additional injury as a result of anoxia. Pain results from a combination of kinins and the effect of lysozymes and pressure from the swelling on nerve endings. Unchecked, the inflammatory response can set off a neural feedback loop and cause hyperalgesia, a phenomenon in which the surrounding area of injury remains painful. Accordingly, there is a great interest in the medical community to develop anti-inflammatory agents.

Many anti-known inflammatory compositions reduce the inflammatory response, but are also immunosuppressive. For example, corticosteroids are potent anti-inflammatory agents, but are associated with T-cell suppression and increased infections. Interleukin-10 (as well as IL-4 and IL-3 to lesser extents) are broadly acting anti-inflammatory agents, but are associated with decreased cell mediated immune functions.

It would be advantageous to provide compositions and methods which provide protection from adverse effects associated with inflammation, preferably without unnecessary immunosuppression. The present invention provides such compositions and methods.

SUMMARY OF THE INVENTION

Compositions and methods for providing protection from adverse effects associated with inflammation are disclosed. The compositions and

methods can suppress plasma concentrations of tissue necrosis factor-alpha (TNF- α) while increasing plasma concentrations of interleukin-6 (IL-6).

The compositions include non-interlinked particles of bioactive glass with a size less than about 20 μm , alone or in combination with an additional anti-inflammatory agent, and optionally include other therapeutic agents. Formulations including the composition and a suitable carrier, preferably for oral, intramuscular, intraperitoneal or intravenous administration, are also disclosed.

The composition can be administered orally, intramuscularly, intraperitoneally or intravenously to provide systemic relief from the adverse effects associated with inflammation, for example, the effects of excess TNF- α .

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a graph showing the number of cells (cells/ml $\times 10^4$) for mice injected intraperitoneally with control and with bioactive glass, as described in Example 1.

Figure 2 is a graph showing the concentration of TNF- α (pg/ml) for mice injected intraperitoneally with control and with bioactive glass, as described in Example 1.

Figure 3 is a graph showing the concentration of IL-6 (pg/ml) for mice injected intraperitoneally with control and with bioactive glass, as described in Example 1.

Figure 4 is a graph showing the concentration of IL-6 (pg/ml) for mice injected intraperitoneally with varying doses of bioactive glass, as

described in Example 2.

Figure 5 is a graph showing the concentration of TNF- α (pg/ml) for mice injected intraperitoneally with bioactive glass and LPS/D-galactosamine, as described in Example 2.

Figure 6 is a graph showing the concentration of TNF- α (pg/ml) for mice injected intraperitoneally with bioactive glass and LPS, as described in Example 2.

Figure 7 is a graph showing the concentration of IL-1 α (pg/ml) for mice injected intraperitoneally with bioactive glass and LPS, as described in Example 2.

Figure 8 is a graph showing the concentration of IL-6 (pg/ml) for mice injected intraperitoneally with bioactive glass and LPS, as described in Example 2.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

Compositions and methods for providing protection from adverse effects associated with inflammation are disclosed. The compositions and methods can suppress plasma concentrations of tissue necrosis factor-alpha (TNF- α) while increasing plasma concentrations of interleukin-6 (IL-6).

The compositions can be administered orally, intramuscularly, intraperitoneally or intravenously to provide systemic relief from the adverse effects associated with inflammation, for example, the effects of excess TNF- α . The bioactive glass is bioactive *in vivo*, and is not pro-inflammatory. It does not cause the recruitment of PMNs and does not stimulate TNF- α secretion. It stimulates IL-1 β release very modestly, and

induces an IL-6 response.

IL-6 is a unique cytokine with pro- and anti-inflammatory properties. It is the primary hepatic acute phase response inducer (a pro-inflammatory effect). It suppresses TNF- α production by macrophages (an anti-inflammatory effect), promotes B-cell proliferation and promotes a Th₂ response, and is mitogenic for some cell types. It has been shown to reduce collagen induced arthritis (Immunology, 95(1):31 (1998)) and to reduce monocyte-mediated TNF production in response to LPS (Exp. Cell Res., 10 215(1):51-56 (1994) and Am. J. Physiol. 267 (4pt.1):L442-446 (1994)).

The compositions include non-interlinked particles of bioactive glass with a size less than about 20 μ m, alone or in combination with an additional anti-inflammatory agents, and optionally include other therapeutic agents. 15 Formulations including the composition and a suitable carrier, preferably for oral or intravenous administration, are also disclosed.

The normal inflammatory effect is generally regarded as being ligand/receptor controlled. Not being bound to any particular theory or 20 mechanism, it is believed that the rate and amount of ions generated by the absorption of the small particles of bioactive glass has an effect on the pro-inflammatory receptors, which is responsible for the suppression of the cytokines involved in the inflammatory process. Further, the surface of the small particles is micro-porous, and may adsorb the cytokines, rendering 25 them inactive in the inflammatory process.

The compositions and methods described herein are advantageous because they reduce the inflammatory response, but are not broadly 30 immunosuppressive. There does not appear to be a typical dose dependency

in terms of particles of bioactive glass and IL-6 response. The response is independent of dose up to extremely high levels of particulate, for example, over XXX mg injection. Further, the bioactive glass particles appear to elicit a specific IL-6 response, transient in nature, lasting on the order of several hours. Pre-exposure to bioactive glass particles results in a significantly attenuated pro-inflammatory IL-6 cytokine response that results in the suppression of a typical inflammatory response caused by injection of an endotoxin after the injection of the bioactive glass particles.

The terms “wound” and “burn,” collectively referred to herein as “injury” have their usual meanings. “Wound” is intended to include wounds caused by surgical procedures. “Normal” is used in the sense it is usually used in the medical arts. “Medical practitioner” means one of ordinary skill in the art wound and burn treatment. Typically, this person is a physician, nurse, dentist, or paramedic.

I. Bioactive Glass

Compositions including non-interlinked particles of bioactive glass with an average diameter of less than about 20 μm , alone or in combination with anti-inflammatory agents, can be used for the methods described herein.

Very small particulate bioactive glass has the property of exerting an anti-inflammatory effect when administered systemically. It appears that the bioactive glass suppresses the production of tissue necrosis factor alpha (TNF- α) and interleukin-6 (IL-6). TNF- α is a powerful pro-inflammatory cytokine that not only participates in the normal inflammatory response, but is also implicated in myocardial dysfunction and cardiomyocyte death in ischemia-reperfusion injury, sepsis, chronic heart failure, viral myocarditis and cardiac allograft rejection, as well as a host of other inflammatory disorders. Accordingly, by suppressing the production of TNF- α , the

compositions reduce the likelihood of these disorders occurring.

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The preferred size range for the bioactive glass, for this embodiment, is such that the particles do not physically obstruct vascular, lymph or pulmonary pathways as the particles pass through the body. As the particles are less than about 20 microns in size, they avoid phagocytosis and uptake by the reticuloendothelial system. This is in stark contrast to small particles of other materials, such as talcum, asbestos, silicone and metal debris, which are known to be strongly pro-inflammatory. The particles are of a suitable size for intravenous administration.

As used herein the terms "bioactive glass" or "biologically active glass" mean an inorganic glass material having an oxide of silicon as its major component and which is capable of bonding with growing tissue when reacted with physiological fluids.

Bioactive glasses are well known to those skilled in the art, and are disclosed, for example, in *An Introduction to Bioceramics*, L. Hench and J. Wilson, eds. World Scientific, New Jersey (1993), the contents of which are hereby incorporated by reference.

The glass preferably includes between 40 and 86% by weight of silicon dioxide oxide (SiO_2), between about 0 and 35% by weight of sodium oxide (Na_2O), between about 4 and 46% by weight calcium oxide (CaO), and between about 1 and 15% by weight phosphorus oxide (P_2O_5). More preferably, the glass includes between 40 and 60% by weight of silicon dioxide oxide (SiO_2), between about 5-30% by weight of sodium oxide (Na_2O), between about 10 and 35% by weight calcium oxide (CaO), and between about 1 and 12% by weight phosphorus oxide (P_2O_5). The oxides can be present as solid solutions or mixed oxides, or as mixtures of oxides.

CaF₂, B₂O₃, Al₂O₃, MgO and K₂O may be included in the composition in addition to silicon, sodium, phosphorus and calcium oxides. The preferred range for B₂O₃ is between 0 and 10% by weight. The preferred range for K₂O is between 0 and 8% by weight. The preferred range for MgO is between 0 and 5% by weight.

The most preferred glass is Bioglass®™ (a trademark of University of Florida), which has a composition including about 45% by weight silicon dioxide, about 24.5% by weight sodium oxide, about 6% by weight phosphorus oxide, and about 24.5% by weight calcium oxide. Another preferred material is hydroxyapatite.

Particulate, non-interlinked bioactive glass is preferred in the present invention. That is, the glass is in the form of small, discrete particles, rather than a fused matrix of particles or a mesh or fabric (woven or non-woven) of glass fibers. Note that under some conditions the discrete particles of the present invention may tend to cling together because of electrostatic or other forces but are still considered to be non-interlinked. The particle size is less than about 20 microns, preferably 10 microns or less, more preferably less than about 5 microns, and ideally, less than about 2 microns.

The glass composition can be prepared in several ways, to provide melt-derived glass, spun fibers of sol-gel derived glass, and sintered glass particles. The sintered particles may be in sol-gel derived, or pre-reacted melt derived form. Sol-gel derived glass is generally prepared by synthesizing an inorganic network by mixing metal alkoxides in solution, followed by hydrolysis, gelation, and low temperature (600-900°C) firing to produce a glass. Melt derived glass is generally prepared by mixing grains of oxides or carbonates, melting and homogenizing the mixtures at high temperatures, typically between about 1250 and 1400°C. The molten glass

can be fritted and milled to produce a powder or casted into steel or graphite molds to make bulk implants.

5 The glass composition is preferably melt-derived. In each preparation, it is preferred to use reagent grade glass, especially since the glass is used to prepare materials which ultimately may be implanted in a human. Other compositions which biodegrade and release the same ions as the bioactive glasses described above can also be used, provided they do not elicit an inflammatory response and that they also elicit enhanced IL-6
10 production.

A. Melt Derived Glass

A melt-derived glass composition can be prepared, for example, by preparing an admixture of the individual metal oxides and other components
15 used to prepare the glass composition, blending the admixture, melting the admixture, and cooling the mixture. The melting temperature is determined in large part by the glass composition, and ranges, for example, from about 900-1500°C, preferably between about 1250 and 1450°C. The melt is preferably mixed, for example, by oxygen bubbling, to ensure a thorough
20 homogenation of the individual components.

The mixture can be cooled, for example, by adding the molten admixture to a suitable liquid, such as deionized water, to produce a glass frit. Porosity can be introduced by grinding the glass into a powder,
25 admixing the powder with a foaming agent, and hot pressing the mixture under vacuum and elevated temperature. The particle size of the glass powder is between about 40 and 70 μm , the vacuum is preferably less than 50 MPa, and the hot pressing is preferably performed at a temperature above 400°C, preferably between about 400 and 500°C. Suitable foaming agents
30 include compounds which evolve carbon dioxide and/or water at elevated

temperatures, for example, metal hydroxides, metal carbonates, and peroxides, such as hydrogen peroxide. Preferred metal carbonates are sodium bicarbonate, sodium carbonate and calcium carbonate. The foaming agents are preferably added in a range of between about 1-5, more preferably 2-3 percent by weight of the glass powder. The preparation of melt-derived porous glass is described, for example, in U.S. Patent No. 5,648,301 to Ducheyne and El Ghannam, the contents of which are hereby incorporated by reference.

B. Sintered Glass Particles

Glass can be sintered using known methodology. In one embodiment, an aqueous slurry of the glass powder and a foaming agent with a suitable binder, such as polyvinyl alcohol, is formed. The slurry is then poured into a mold, allowed to dry, and sintered at high temperatures. These temperature may range, depending on the glass composition and foaming agent used, between about 500 and 1000°C, more preferably between about 600 and 800°C.

C. Spun Fibers of Sol-gel Derived Glass

It is known in the art to control the heat treatment cycle of glass gels to control the pores and interpores of the material to create a porous glass material. Suitable pore diameters are between 20 and 180Å, suitable pore volumes are between 40 and 52 cc/g, and suitable surface areas are between 75 and 350 m²/g. Since a pore diameter larger than 0.1 microns is difficult to achieve using this method, the sintering and foaming processes described herein are generally more preferred.

D. Leaching of the Porous Material

To aid in preparing glass compositions with high porosity, the glass composition can include a material which can be preferably leached out of

the glass composition, and, in doing so, provide the composition with high porosity. For example, minute particles of a material capable of being dissolved in a suitable solvent, acid, or base can be mixed with or melted into the glass, and subsequently leached out. The resulting voids have roughly the same size as the particle that was leached out. In the case of a material which is part of a melt-derived glass composition, the size of the pores and degree of porosity depends on the amount of added material relative to the amount of glass. For example, if the leached material constituted about 80% of the glass, then the glass would be approximately 80% porous when the material was leached out. When leaching the glass composition, care should be taken not to leach out those components which add to the bioactivity of the glass, i.e., the calcium and phosphorus oxides.

II. Formulations including Bioactive Glass

The bioactive glass particulates are preferably administered in a formulation that includes an acceptable carrier for the mode of administration. Suitable pharmaceutically acceptable carriers are known to those of skill in the art. The formulations can optionally include other therapeutically active ingredients, such as antibiotics, antivirals, healing promotion agents, anti-inflammatory agents, immunosuppressants, growth factors, anti-metabolites, cell adhesion molecules (CAMs), antibodies, vascularizing agents, anti-coagulants, and anesthetics/analgesics.

The carrier must be pharmaceutically acceptable in the sense of being compatible with the other ingredients of the formulation and not deleterious to the recipient thereof. The formulations can include carriers suitable for oral, rectal, topical or parenteral (including subcutaneous, intramuscular and intravenous) administration. Preferred carriers are those suitable for oral or parenteral administration.

Formulations suitable for parenteral administration conveniently include a sterile aqueous preparation of the bioactive glass. Such formulations may conveniently contain distilled water, 5% dextrose in distilled water or saline. The formulations may be in the form of a powdered formulation including the bioactive glass particulates and salt, dextrose, buffers and the like, which can be dissolved in water immediately prior to administration.

For enteral administration, the bioactive glass particulates can be incorporated into an inert carrier in discrete units such as capsules, cachets, tablets or lozenges, each containing a predetermined amount of the particles; as a powder or granules; or a suspension or solution in an aqueous liquid or non-aqueous liquid, *e.g.*, a syrup, an elixir, an emulsion or a draught. Suitable carriers may be starches or sugars and include lubricants, flavorings, binders, and other materials of the same nature.

A tablet may be made by compression or molding, optionally with one or more accessory ingredients. Compressed tablets may be prepared by compressing in a suitable machine the active compound in a free-flowing form, *e.g.*, a powder or granules, optionally mixed with accessory ingredients, *e.g.*, binders, lubricants, inert diluents, surface active or dispersing agents. Molded tablets may be made by molding in a suitable machine, a mixture of the powdered active compound with any suitable carrier.

A syrup or suspension may be made by adding the active compound to a concentrated, aqueous solution of a sugar, *e.g.*, sucrose, to which may also be added any accessory ingredients. Such accessory ingredients may include flavoring, an agent to retard crystallization of the sugar or an agent to increase the solubility of any other ingredient, *e.g.*, as a polyhydric alcohol,

for example, glycerol or sorbitol.

In addition to the aforementioned ingredients, the formulations may further include one or more optional accessory ingredient(s) utilized in the art of pharmaceutical formulations, e.g., diluents, buffers, flavoring agents, binders, surface active agents, thickeners, lubricants, suspending agents, preservatives (including antioxidants) and the like.

Other Therapeutic Agents

In addition to bioactive glass particles, the formulations can include other therapeutic agents such as antibiotics, antivirals, healing promotion agents, anti-inflammatory agents, immunosuppressants, growth factors, anti-metabolites, cell adhesion molecules (CAMs), bone morphogenic proteins (BMPs), vascularizing agents, anti-coagulants, and topical anesthetics/analgesics.

The antibiotics can be topical antibiotics suitable for skin treatment. Examples of such antibiotics include but are not limited to: chloramphenicol, chlortetracycline, clyndamycin, clioquinol, erythromycin, framycetin, gramicidin, fusidic acid, gentamicin, mafenide, mupirocin, neomycin, polymyxin B, bacitracin, silver sulfadiazine, tetracycline and chlortetracycline.

Suitable antivirals include topical antivirals, such as acyclovir and gancyclovir. Suitable anti-inflammatory agents include corticosteroids, hydrocortisone and nonsteroidal antinflammatory drugs. Suitable growth factors include basic fibroblast growth factor (bFGF), epithelial growth factor (EGF), transforming growth factors α and β (TGF α and β), platelet-derived growth factor (PDGF), and vascular endothelial growth

factor/vascular permeability factor (VEGF/VPF)). Suitable topical anesthetics include benzocaine and lidocaine.

An additional anti-inflammatory agent which can be used in combination with the bioactive glass particles is β -glucan. β -glucan is a polysaccharide with anti-inflammatory properties, and, like the bioactive glass particles described herein, increases IL-6 release and decreases TNF- α and IFN γ release.

10 **III. Methods of Reducing Inflammation**

The compositions can be used to prophylactically or therapeutically to reduce inflammation in a patient. Overly acute or chronic inflammation can result in various disease states in a patient, for example, arthritis and tendonitis, pulmonary disorders such as asthma and emphysema, and post-surgical (peritoneal) adhesions.

The particles of bioactive glass are believed to modify the inflammatory response in the local microenvironment by altering the synthetic properties of resident macrophages and other recruited inflammatory cells. The particles induce macrophage tolerance by eliciting a cytokine response (IL-6 based) that acts in an autocrine and paracrine fashion.

Particles of bioactive glass can be delivered by intravenous, intramuscular, or intraperitoneal injection to provide systemic (and possibly local when administered intramuscularly or intraperitoneally) anti-inflammatory effects. These effects can be therapeutic and/or prophylactic. For example, systemic delivery of bioactive glass can be effective in reducing the onset of inflammation brought on by external challenge.

Particles of bioactive glass can also be administered orally.

Systemic administration of the bioactive glass particles can lower production of TNF- α and also increase production of IL-6. The effect of this is to modify the pro-inflammatory response, and will be useful in treating or preventing sepsis, systemic inflammatory response syndromes, adult
5 respiratory distress syndrome, ocular injury, surgical wound healing and adhesion formation, and delayed cutaneous wound healing. Other disorders which are, at least in part, attributable to an exaggerated inflammatory response, such as ARDS, pancreatitis, viral hepatitis, hemorrhagic shock, ischemia/reperfusion injury, peritoneal adhesions, and chronic inflammation,
10 such as delayed wound healing and rheumatoid arthritis, can be treated with systemic administration of bioactive glass particles.

In one embodiment, the particles are administered locally, for example, by inhalation, by spraying (in the form of an aerosol) or by mixing
15 the particles with a gel (for example, a biocompatible hydrogel), creme or other aqueous or non-aqueous carrier, and applying or injecting the composition subcutaneously at a site at which surgery is to be performed at a later time. The compositions can advantageously include an anesthetic. The presence of the particles at the proposed surgical site can lead to decreased
20 inflammation resulting from the surgery.

The present invention will be more clearly understood with reference to the following non-limiting examples.

25 **EXAMPLES**

Example 1: Intraperitoneal Administration of Bioactive Glass to Mice

Ten mice were injected intraperitoneally with 25 mgs of bioactive glass (45S5) with a particle size less than about 20 μ m in a total volume of 1
30 ml (0.5 ml fetal calf serum and 0.5 ml phosphate-buffered saline) with a

result pH of 9.6. An additional ten mice received the carrier (0.5 ml fetal calf serum and 0.5 ml phosphate-buffered saline) with the pH unadjusted. Two hours later, the mice were euthanized and the peritoneal contents were washed with 3 ml of physiologic saline. Peritoneal white cell count was performed with a hemocytometer, and peritoneal TNF- α and IL-6 determined by ELISA.

The procedures were performed without incident. The animals appeared grossly normal for the two hour post-injection period. At sacrifice, peritoneal white cell count in mice receiving the bioactive glass particles was not different from peritoneal wash fluid from mice receiving only the carrier. The proinflammatory cytokine TNF- α was not detected in any of the samples. Peritoneal IL-6 concentrations, however, were increased 25 fold from approximately 80 pgs/ml in the carrier-treated mice to over 2,000 pgs/ml in the bioactive glass-treated mice. The cell count, TNF- α concentration and IL-6 concentration from the peritoneal washings for the bioactive glass-treated mice and the control mice are shown in Figures 1-3.

In conclusion, the bioactive glass is bioactive when administered intraperitoneally. The bioactive glass is not acting as a classical irritant. A chemical irritation or a direct inflammatory agent like endotoxin or TNF- α would have resulted in a significant and rapid neutrophil influx, which was not observed. The bioactive glass was not directly pro-inflammatory, since no TNF- α response was elicited. A significant IL-6 response was observed, but IL-6 has both pro-inflammatory (hepatic acute phase induction) and anti-inflammatory properties (delays apoptosis, suppresses macrophage TNF and IL-1 production). Since no inflammatory cell influx was observed, the enhanced IL-6 production must have been secondary to resident macrophages and fibroblasts. The results are consistent with bioactive glass-

mediated stimulation of resident cell IL-6 synthesis, which represents a new anti-inflammatory property.

Example 2: Anti-Inflammatory Effect of Bioactive Glass in a Mouse Endotoxycosis Model

Bioactive glass particles with a particle size of 5 μm were injected to C57B1/6 mice. The mice were injected ip with 3, 6, 12, 25 and 50 mg of bioactive glass particles or buffer. The animals were sacrificed and TNF- α , IL-1 α and IL-6 levels were determined in the peritoneal lavage fluid at 2 hours (Figure 4). In a second study, mice were injected with 12 mg of bioactive glass particles or buffer ip and then challenged ip with 1 μg of LPS and 8 mg of D-galactosamine (Dgal) 6 hours after the bioactive glass particles were administered. Blood samples were taken 90 minutes after Dgal/LPS injection for determination of TNF- α . Lastly, a sublethal dose of LPS (100 ng) was given ip 2 hours after an ip injection of 3 mg of 5 μm particles of bioactive glass. The mice were bled, sacrificed and lavaged 2 hours after the LPS administration.

Results: In the first study, all doses of bioactive glass were found to induce a significant IL-6 response (range 4,584-23,112 pg/ml bioactive glass, $1,017 \pm 445$ pg/ml \pm SEM buffer); however, the bioactive glass did not induce TNF- α or IL-1 α production in the peritoneal lavage fluid. In the second study, mice that received bioactive glass particles prior to Dgal/LPS had significantly lower plasma TNF- α than did controls (bioactive glass: $13,047 \pm 4,126$ pg/ml, control: $34,813 \pm 4,902$ pg/ml, $p < 0.0038$) (Figure 5). Finally, peritoneal IL-6 and TNF- α were reduced in response to LPS by pretreatment with bioactive glass (IL-6 $24,452 \pm 6,673$ pg/ml and control: $53,330 \pm 2,586$ pg/ml, bioactive glass: TNF- α - 26 ± 5 pg/ml and control: 91 ± 49 pg/ml) (Figures 6-8).

Conclusion: Bioactive glass particles are a bioactive substance that when administered alone appear to elicit a significant initial IL-6 response without concurrent expression of TNF- α or IL-1 α .

We claim:

1. A composition comprising particles of bioactive glass with a particle size less than about 20 μm in diameter and a suitable carrier for oral, intramuscular, intraperitoneal or intravenous administration.

5

2. The composition of claim 1, wherein the carrier is suitable for intravenous administration.

3. The composition of claim 1, wherein the carrier is suitable for intramuscular or intraperitoneal administration.

10

4. The composition of claim 1, additional comprising one or more therapeutic agents.

15

5. The composition of claim 4, wherein one or more therapeutic agents are selected from the group consisting of healing promotion agents, growth factors, anti-inflammatory agents, and anesthetics.

20

6. The composition of claim 1 wherein the glass includes between about 40 and 86 percent by weight of SiO_2 , between about 0 and 30 percent by weight of Na_2O , between about 4 and 46 percent by weight of CaO and between about 1 and 15 percent by weight of P_2O_5 .

25

7. The composition of claim 1, wherein the bioactive glass has a particle size range less than about 2 microns.

30

8. A method for systemically minimizing the production of $\text{TNF-}\alpha$ caused by an inflammatory response in a patient comprising orally or intravenously administering an effective $\text{TNF-}\alpha$ lowering amount of bioactive glass particles with a size less than about 20 μm to the patient.

9. The method of claim 8 wherein the systemic minimization of TNF- α is used prophylactically or therapeutically to prevent or treat a disease state selected from the group consisting of peritoneal adhesions, septic shock, cardiac allograft rejection, systemic inflammatory response syndromes, adult respiratory distress syndrome, ocular injury, delayed cutaneous wound healing, ARDS, pancreatitis, viral hepatitis, hemorrhagic shock, ischemia, reperfusion injury, and rheumatoid arthritis.

10. A method for systemically increasing IL-6 levels in a patient, comprising administering to the patient an effective, IL-6 increasing amount of bioactive glass particles with a size less than about 20 μm .

11. A composition comprising particles of a material with a particle size less than about 20 μm which biodegrades, produces elevated serum concentrations of calcium and phosphorous ions, does not cause elevated plasma TNF- α concentrations, and does cause elevated plasma IL-6 concentrations, in combination with a suitable carrier for oral, intramuscular, intraperitoneal or intravenous administration.

20

Compositions and methods for systemically minimizing the inflammatory effects of TNF- α are disclosed. The compositions include particles of bioactive glass with a particle size less than about 20 μm , alone or in combination with anti-inflammatory agents and other therapeutic agents. The compositions can include an appropriate carrier for oral, intramuscular, intraperitoneal or intravenous administration. When administered to a patient, the particles bring about elevated levels of IL-6 and do not cause elevated levels of TNF- α .

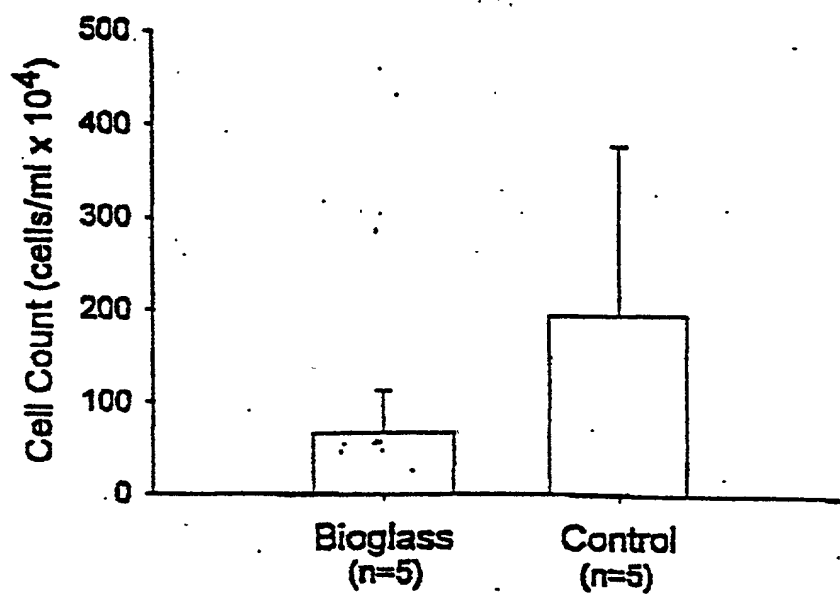


Fig. 1

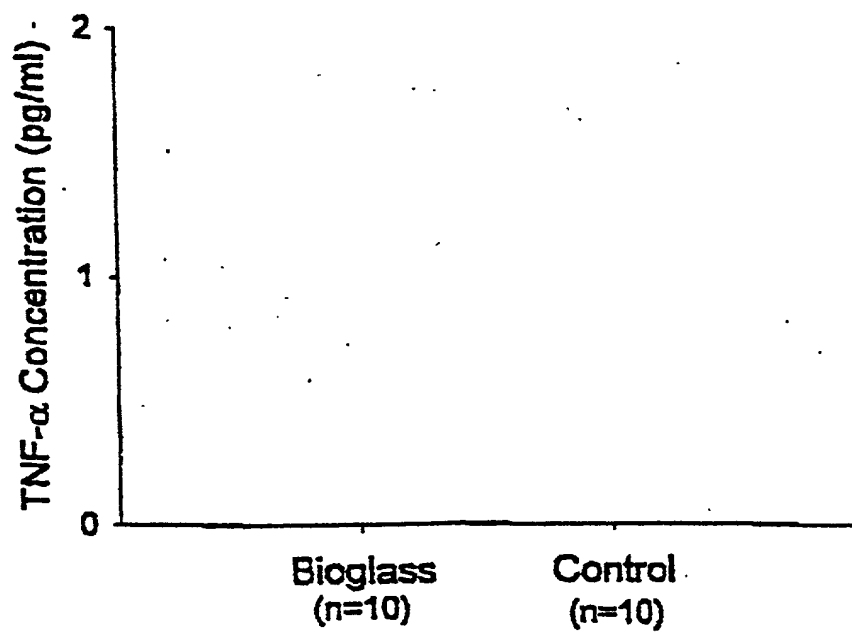


Fig. 2

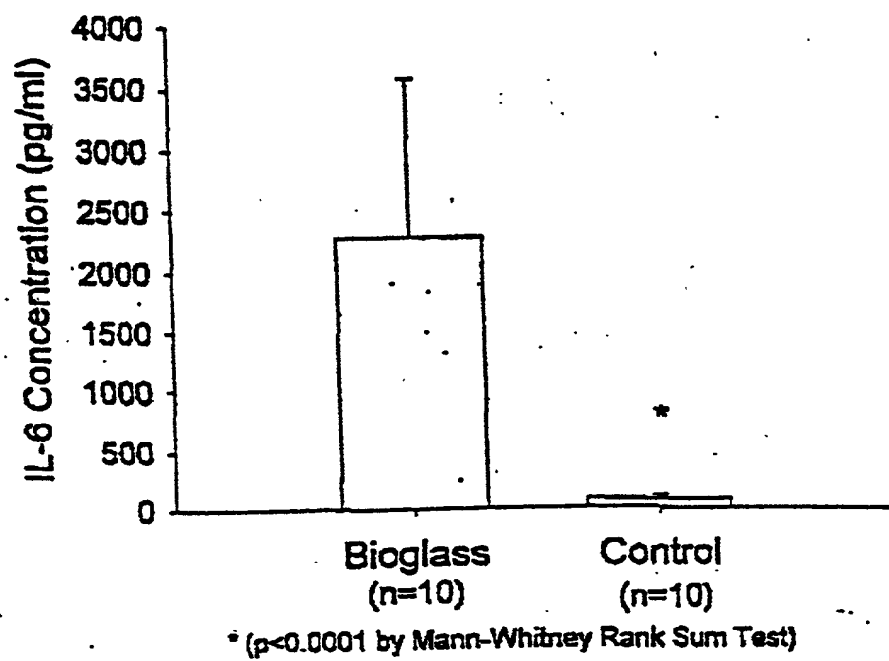


Fig. 3

IL-6 Levels in Peritoneal Lavage Fluid Following Varying Doses of ip Bioglass

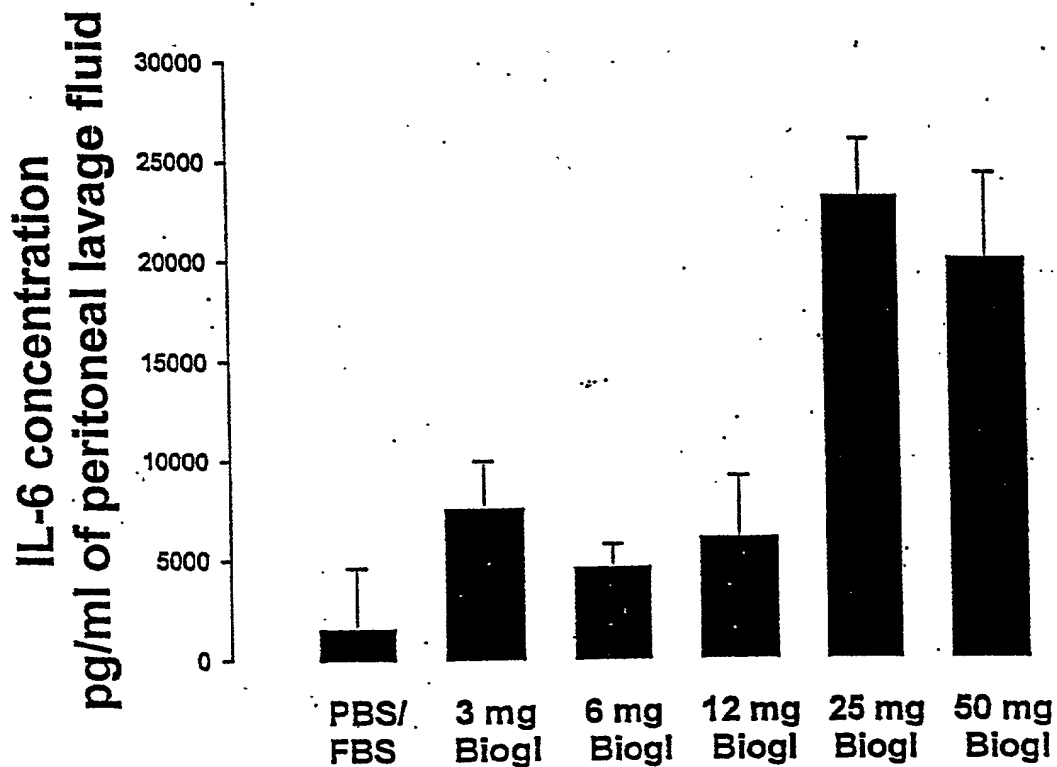


Fig. 4

TNF Concentration in Mice Injected with Bioglass and LPS/D-galactosamine

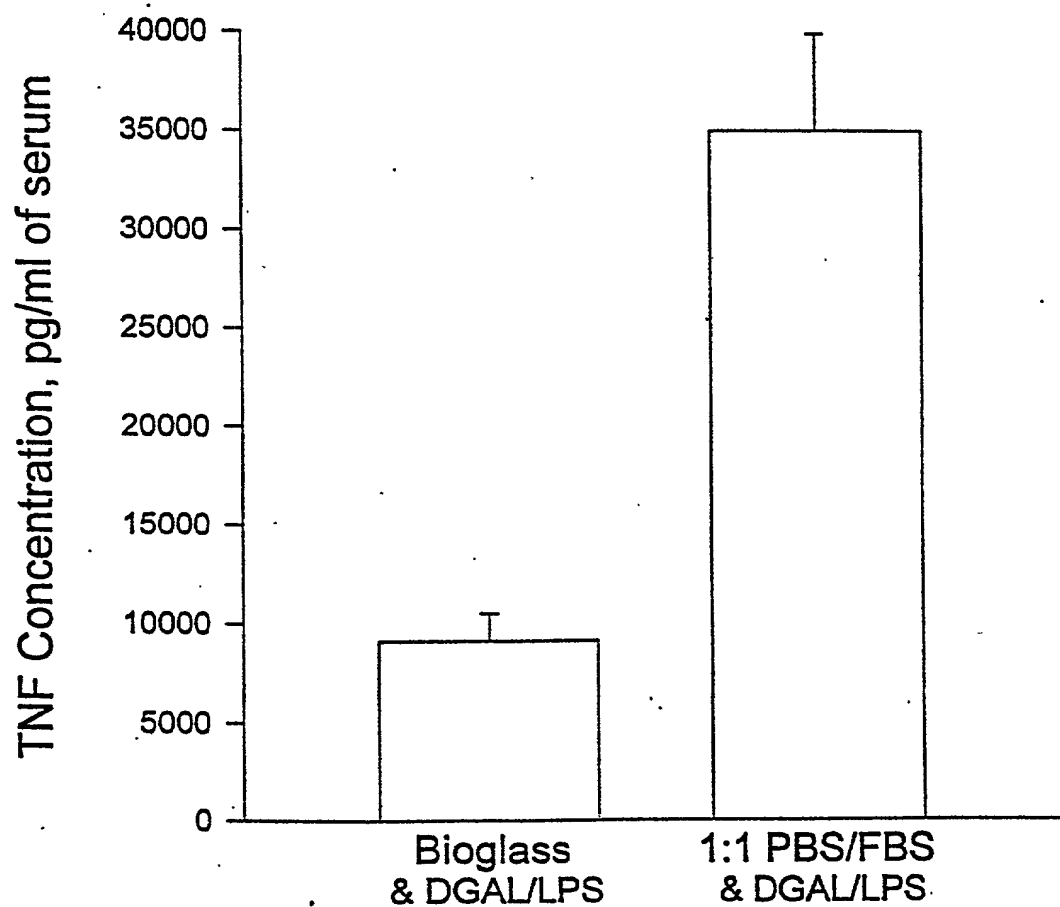


Fig. 5

TNF α Concentration in Peritoneal Lavage Fluid Following ip Injection of Bioglass and LPS

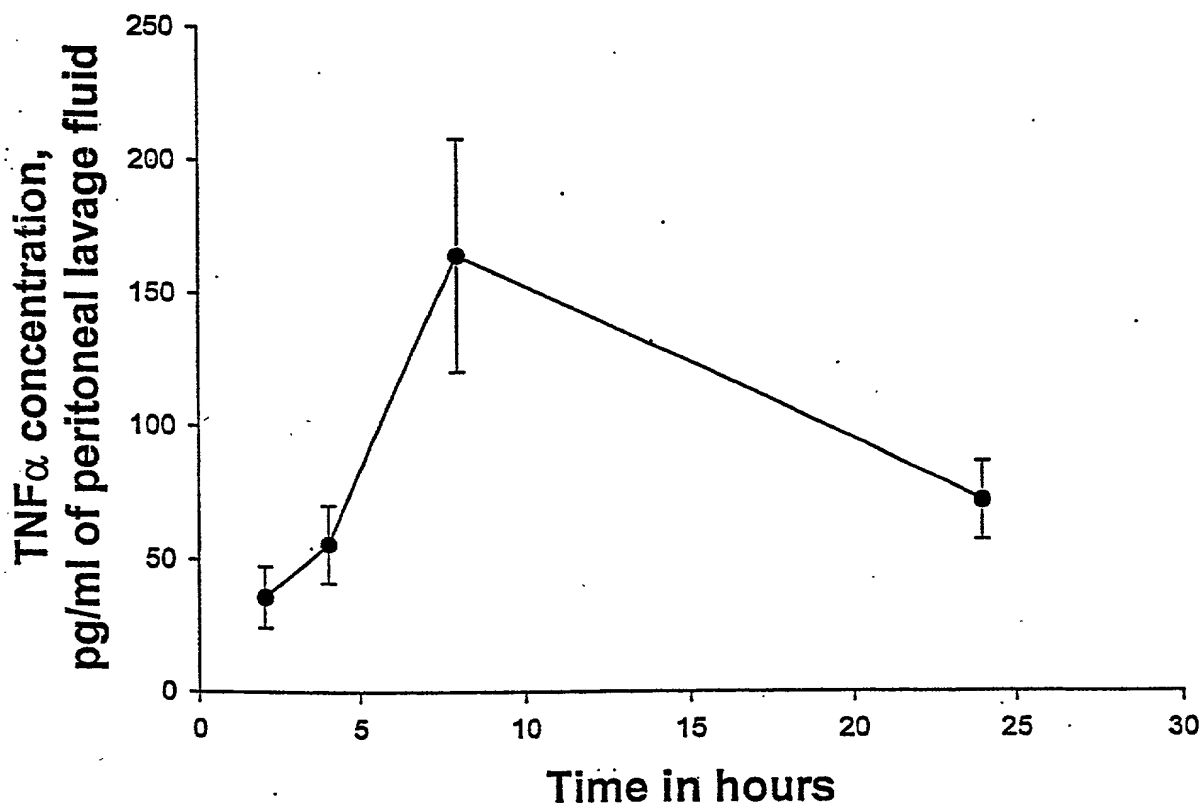


Fig. 6

Concentration of IL1- α in Peritoneal Lavage Fluid of Mice Given Bioglass and LPS

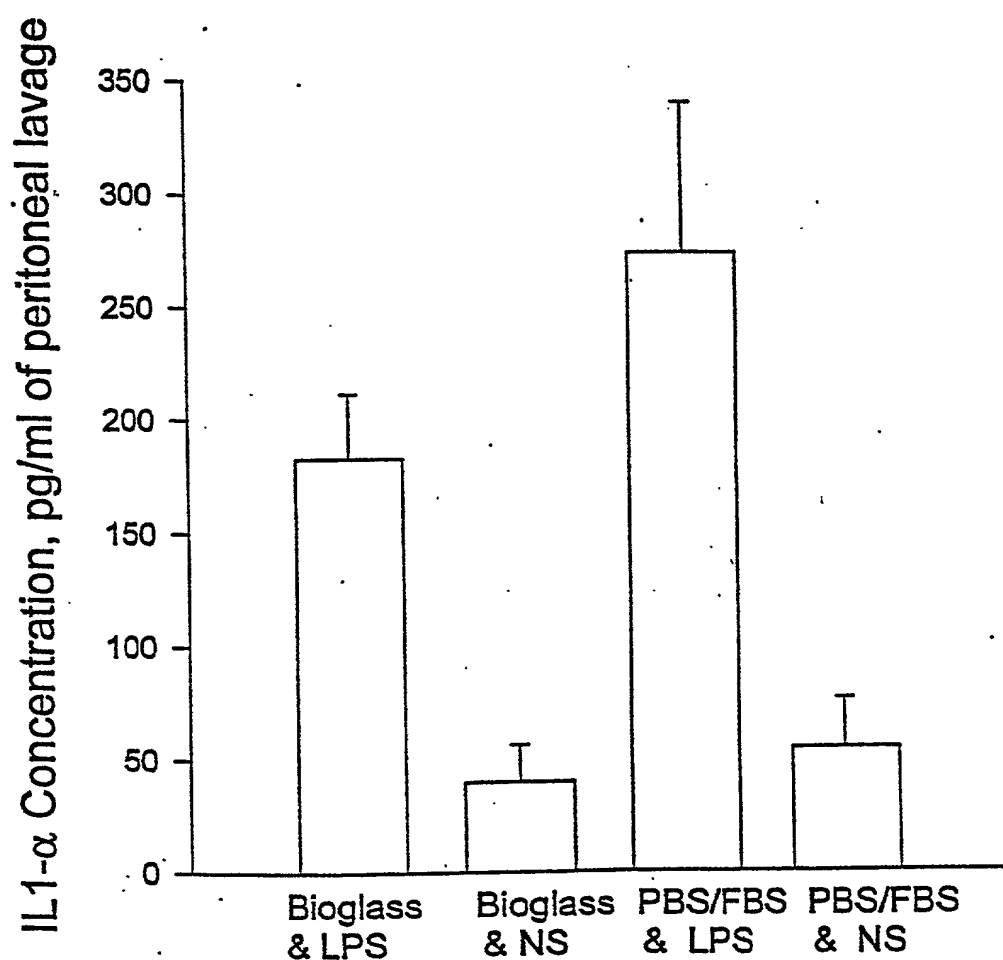


Fig. 7

IL-6 Concentration in Mice Injected with Bioglass and LPS

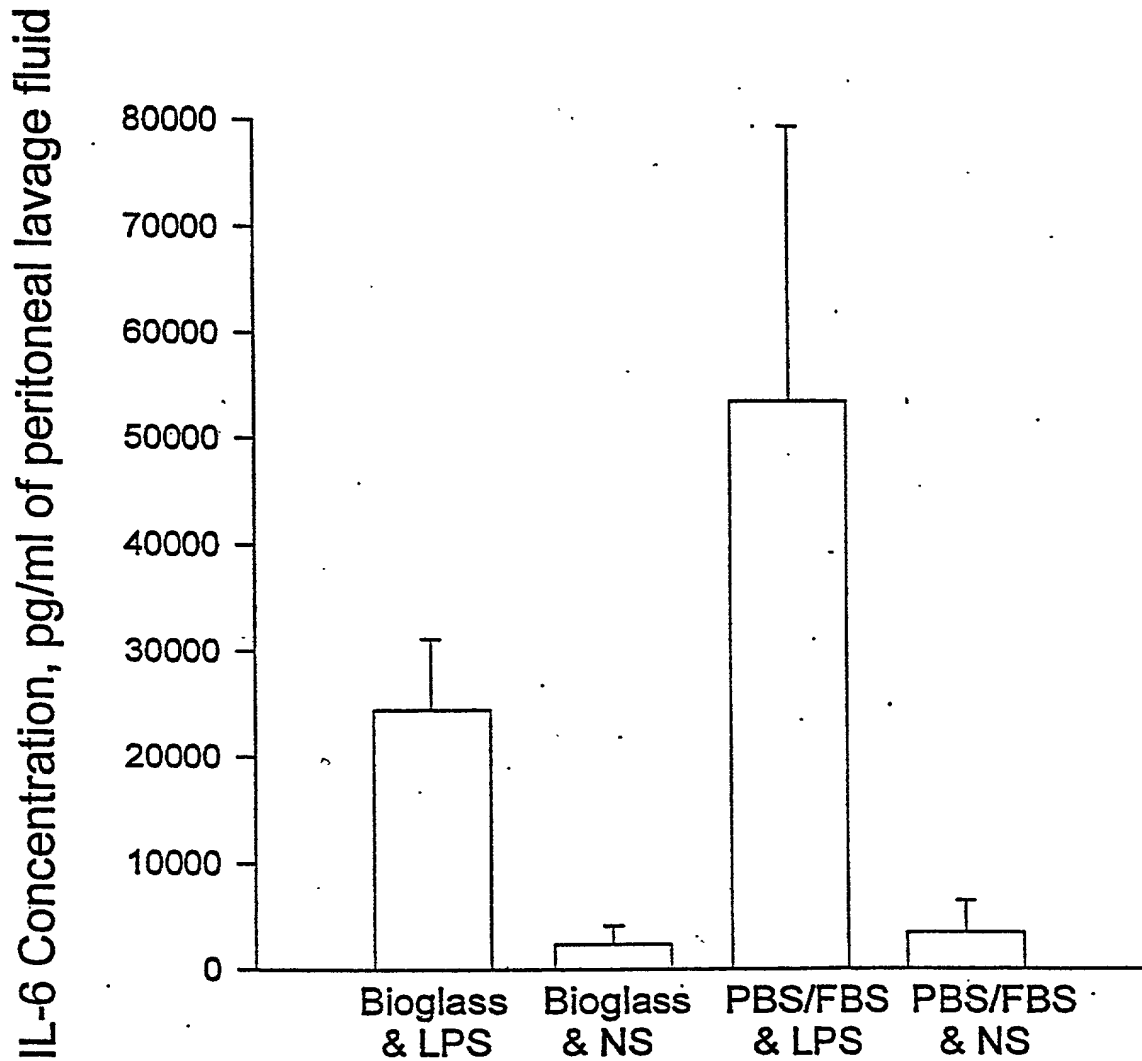


Fig. 8

**COMBINED DECLARATION AND POWER OF ATTORNEY
FOR UTILITY PATENT APPLICATION**

Attorney's Docket No.

028870-178

As a below-named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name;

I BELIEVE I AM THE ORIGINAL, FIRST AND SOLE INVENTOR (if only one name is listed below) OR AN ORIGINAL, FIRST AND JOINT INVENTOR (if more than one name is listed below) OF THE SUBJECT MATTER WHICH IS CLAIMED AND FOR WHICH A PATENT IS SOUGHT ON THE INVENTION ENTITLED:

Anti-Inflammatory Bioactive Glass Particulates

the specification of which

(check one)

☒ is attached hereto;

☐ was filed on _____ as

Application No. _____

and was amended on _____;
(if applicable)

I HAVE REVIEWED AND UNDERSTAND THE CONTENTS OF THE ABOVE-IDENTIFIED SPECIFICATION, INCLUDING THE CLAIMS, AS AMENDED BY ANY AMENDMENT REFERRED TO ABOVE;

I ACKNOWLEDGE THE DUTY TO DISCLOSE TO THE OFFICE ALL INFORMATION KNOWN TO ME TO BE MATERIAL TO PATENTABILITY AS DEFINED IN TITLE 37, CODE OF FEDERAL REGULATIONS, Sec. 1.56 (as amended effective March 16, 1992);

I do not know and do not believe the said invention was ever known or used in the United States of America before my or our invention thereof, or patented or described in any printed publication in any country before my or our invention thereof or more than one year prior to said application; that said invention was not in public use or on sale in the United States of America more than one year prior to said application; that said invention has not been patented or made the subject of an inventor's certificate issued before the date of said application in any country foreign to the United States of America on any application filed by me or my legal representatives or assigns more than twelve months prior to said application;

I hereby claim foreign priority benefits under Title 35, United States Code Sec. 119 and/or Sec. 365 of any foreign application(s) for patent or inventor's certificate as indicated below and have also identified below any foreign application for patent or inventor's certificate on this invention having a filing date before that of the application(s) on which priority is claimed:

COMBINED DECLARATION AND POWER OF ATTORNEYAttorney's Docket No.
028870-178

COUNTRY/INTERNATIONAL	APPLICATION NUMBER	DATE OF FILING (day, month, year)	PRIORITY CLAIMED
U.S.	60/131,529	29.04.99	YES <input checked="" type="checkbox"/> NO <input type="checkbox"/>
			YES <input type="checkbox"/> NO <input type="checkbox"/>

I hereby appoint the following attorneys and agent(s) to prosecute said application and to transact all business in the Patent and Trademark Office connected therewith and to file, prosecute and to transact all business in connection with international applications directed to said invention:

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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COMBINED DECLARATION AND POWER OF ATTORNEYAttorney's Docket No.
028870-178

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